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- L3 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN AN 2009:1224075 CAPLUS < LOGINID::20091101>>
- TI Increased expression of regulatory \*\*\*Tr1\*\*\* cells in recurrent hepatitis Cafter liver transplantation
- AU Carpentier, A.; Conti, F.; Stenard, F.; Aoudiehane, L.; Miroux, C.; Podevin, P.; Morales, O.; Chouzenoux, S.; Scatton, O.; Groux, H.; Auriault, C.; Calmus, Y.; Pancre, V.; Delhem, N. CS Laboratoire de Biologie Cellulaire, Universite Rene Descartes.
- Paris, Fr. SO American Journal of Transplantation (2009), 9(9), 2102-2112 CODEN: AJTMBR; ISSN: 1600-6135
- PB Wiley-Blackwell
- DT Journal
- LA English
- AB Immune response failure during HCV infection has been assocd, with the activity of regulatory T cells. Hepatitis C-related cirrhosis is the main reason for liver transplantation. However, 80% of transplanted patients present an accelerated recurrence of the disease. This study assessed the involvement of regulatory T-cell subsets (CD4+CD25+ cells: 'Treg' and 'CD49h\*\*\* +CD18+ cells: 'T regulatory-1' cells), in the recurrence of HCV after liver transplantation, using transcriptomic anal., ELISA assays on serum samples and immunohistochem, on liver biopsies from liver recipients 1 and 5 years after transplantation. Three groups of patients were included: stable HCV-neg, recipients and those with mild and severe hepatitis C recurrence. At 5 years, Treg markers were overexpressed in all HCV+ recipients. By contrast, """Tr1"" markers were only overexpressed in patients with severe recurrence. At 1 yr, a trend toward the overexpression of \*\*\* Tr1\*\*\* was noted in patients evolving toward severe recurrence. IL-10 prodn., a characteristic of the \*\*\*Tr1\*\*\* subset, was enhanced in severe recurrence at both 1 and 5 years. These results suggest that \*\*\*Tr1\*\*\* are enhanced during severe HCV recurrence after liver transplantation and could be predictive of HCV recurrence. High levels of IL-10 at 1 yr could be predictive of severe recurrence, and high IL-10 producers might warrant more intensive management. RE ONT 41 THERE ARE 41 CITED REFERENCES AVAILABLE
- FORMAT L3 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ALL CITATIONS AVAILABLE IN THE RE

- AN 2005:618 CAPLUS << LOGINID::20091101>> DN 142:73419 TI Method for identification of \*\*\*TR1\*\*\* regulator
- lymphocytes by the presence and the expression of specific molecules, and diagnostic and therapeutic applications IN Grouy Herve
- PA Txcell Fr

FOR THIS RECORD

- SO Fr. Demande, 89 pp. CODEN: FRXXBL
- IA French
- DT Patent
- FAN. ONT 1 PATENT NO. KIND DATE APPLICATION DATE -----

PI FB 2856700 A1 20041231 FR 2003-7601 20030624 FR 2856700 B1 20070608 AU 2004251462 A1 20050106 AU 2004-251462 20040624 CA 2529941 20050106 CA 2004-2529941 20040624 WO 2005000344 A2 20050106 WO 2004-FR1583 A3 20050609 W: AE AG. 20040624 WO 2005000344 AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH. CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC. LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA,

NI. NO. NZ. OM. PG. PH. PL. PT. RO. RU. SC. SD. SE. SG. SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN. YU. ZA, ZM. ZW RW; BW, GH, GM, KE, LS, MW, MZ, NA, SD SL SZ TZ UG ZM ZW AM AZ, BY, KG, KZ, MD. RU. TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK. EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN TD TG EP 1660118 A2 20060531 EP 2004-767439 20040624 FP 1660118 B1 20080409 R: AT, BE, CH, DE DK ES, FR GB, GR IT, LI, LU, NL, SE, MC, PT, IE.SI. FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK JP 2007520197 20070726 JP 2006-516319 20040624 AT 391513 20080415 AT 2004-767439 20040624 ES 2305836

T3 20081101 ES 2004-767439 20040624 A 20030624 WO 2004-FR1583 PRAI FR 2003-7601

W 20040624

AB The invention discloses a method for the identification of \*\*\* TR1\*\*\* regulator lymphocytes in a biol. sample based on the detn, of the simultaneous presence of CD4, CD18 and/or \*\*\* CD11a\*\*\* , \*\*\* CD49b\*\*\* and, if necessary, by the description of a overexpression of genes coding for the mols. CD4. PSGL-1. PECAM-1 and alpha V/.beta.3. The invention also discloses a method for quantification and a method for diagnosis of autoimmune or inflammatory diseases based on the aforementioned process of identification. The invention further discloses a method for enrichment of \*\*\*TR1\*\*\* regulator lymphocytes based on the detn. of the simultaneous presence of these mols. The invention also discloses the use of a compn. so enriched for the treatment by an autoimmune or inflammatory disease, in particular Orohn's disease.

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